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## Dulaglutide

Cat. No.:	HY-P0120
CAS No.:	923950-08-7
Target:	GCGR
Pathway:	GPCR/G Protein
Storage:	Store at 4°C, do not freeze

# Dulaglutide

### BIOLOGICAL ACTIVITY

<b>Description</b>	Dulaglutide (LY2189265) is a glucagon-like peptide-1 (GLP-1) receptor agonist. Dulaglutide can be used for the research of type 2 diabetes (T2D) <sup>[1][2]</sup> .									
<b>IC<sub>50</sub> &amp; Target</b>	GLP-1 receptor <sup>[1]</sup>									
<b>In Vitro</b>	<p>Dulaglutide (50 nM and 100 nM; 24 h) ameliorates ox-LDL-induced oxidative stress and suppresses ox-LDL-induced mitochondrial dysfunction in human aortic endothelial cells (HAECs)<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Human aortic endothelial cells (HAECs)</td> </tr> <tr> <td>Concentration:</td> <td>50 nM, 100 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Suppressed ox-LDL-induced reduction of cell viability and release of lactate dehydrogenase (LDH).</td> </tr> </table>		Cell Line:	Human aortic endothelial cells (HAECs)	Concentration:	50 nM, 100 nM	Incubation Time:	24 hours	Result:	Suppressed ox-LDL-induced reduction of cell viability and release of lactate dehydrogenase (LDH).
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<b>In Vivo</b>	<p>Dulaglutide (0, 0.05, 0.5, 1.5, or 5 mg/kg; s.c.; twice week, for 93 weeks) increases the incidence of thyroid C-cell hyperplasia and neoplasia in the rat carcinogenicity study<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Rats and Transgenic mice<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>0, 0.05, 0.5, 1.5, or 5 mg/kg; 0, 0.3, 1, or 3 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>SC, twice week, for 93 weeks; SC, twice week, for 26 weeks</td> </tr> <tr> <td>Result:</td> <td>Statistically increased diffuse C-cell hyperplasia and adenomas at 0.5 mg/kg. Decreased Systemic exposures over time in mice.</td> </tr> </table>		Animal Model:	Rats and Transgenic mice <sup>[1]</sup>	Dosage:	0, 0.05, 0.5, 1.5, or 5 mg/kg; 0, 0.3, 1, or 3 mg/kg	Administration:	SC, twice week, for 93 weeks; SC, twice week, for 26 weeks	Result:	Statistically increased diffuse C-cell hyperplasia and adenomas at 0.5 mg/kg. Decreased Systemic exposures over time in mice.
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- J Clin Endocrinol Metab. 2022 Feb 9.

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## REFERENCES

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- [1]. Hertzel C Gerstein, et al. Dulaglutide and cardiovascular outcomes in type 2 diabetes (REWIND): a double-blind, randomised placebo-controlled trial. Lancet
- [2]. Chang W, et al. Glucagon-like peptide-1 receptor agonist dulaglutide prevents ox-LDL-induced adhesion of monocytes to human endothelial cells: An implication in the treatment of atherosclerosis. Mol Immunol. 2019 Dec;116:73-79.
- [3]. Byrd RA, et al. Chronic Toxicity and Carcinogenicity Studies of the Long-Acting GLP-1 Receptor AgonistDulaglutide in Rodents. Endocrinology. 2015 Jul;156(7):2417-28.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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